§ 5.302 Detention of meat, poultry, eggs, and related products.

The Regional Food and Drug Directors and District Directors are authorized to perform and to designate other officials to perform all of the functions of the Commissioner of Food and Drugs under:

(a) Section 409(b) of the Federal Meat Inspection Act (21 U.S.C. 679(b)), that relates to the detention of any carcass, part thereof, meat, or meat product of cattle, sheep, swine, goats, or equines.

(b) Section 24(b) of the Poultry Products Inspection Act (21 U.S.C. 467f (b)) that relates to the detention of any poultry carcass, part thereof, or poultry product.

(c) The Egg Products Inspection Act (21 U.S.C. 1031 *et seq.*).

§ 5.303 Establishing standards and approving accrediting bodies under the National Laboratory Accreditation Program.

The Director, Deputy Director, and Director of Regulations and Policy, Center for Food Safety and Applied Nutrition, are authorized to perform all the functions of the Commissioner of Food and Drugs under sections 1322(b) and (c) of the Food, Agriculture, Conservation, and Trade Act of 1990 (the National Laboratory Accreditation Program) (7 U.S.C. 138a), as amended hereafter, which relate to setting standards for the National Laboratory Accreditation Program and approving State agencies or private, nonprofit entities as accrediting bodies to implement certification and quality assurance programs in accordance with the requirements of these sections. The delegation excludes the authority to submit reports to the Congress. These officials may not further redelegate this authority.

§ 5.304 Approval of schools providing food-processing instruction.

(a) The following officials are authorized to perform all the functions of the Commissioner of Food and Drugs (Commissioner) under § 113.10 of this chapter regarding the approval of schools giving instruction in retort operations, processing systems operations, aseptic processing and packaging system operations, and container closure inspections:

(1) The Director and Deputy Director, Center for Food Safety and Applied Nutrition (CFSAN).

(2) The Director of Regulations and Policy, CFSAN.

(3) The Director, Office of Plant and Dairy Foods and Beverages, CFSAN.

(b) These officials may not further redelegate this authority.

Subpart F—Medical Devices and Radiological Health; Redelegations of Authority

§ 5.400 Issuance of Federal Register documents to recognize or to withdraw recognition of a standard to meet premarket submission requirements.

(a) The Director and Deputy Directors for Science and for Regulations and Policy, Center for Devices and Radiological Health; and the Director and Deputy Directors, Center for Biologics Evaluation and Research, are authorized to issue FEDERAL REGISTER documents under section 514(c) of the Federal Food, Drug and Cosmetic Act (the act) (21 U.S.C. 360d(c)) recognizing or withdrawing recognition of a standard for which a person may submit a declaration of conformity in order to meet a premarket submission requirement.

(b) These officials may not further redelegate this authority.

§ 5.401 Issuance of Federal Register documents pertaining to exemptions from premarket notification.

(a) The Director and Deputy Directors for Science and for Regulations and Policy, Center for Devices and Radiological Health; and the Directors and Deputy Directors, Center for Biologics Evaluation and Research, are authorized to make determinations and issue FEDERAL REGISTER notices and rules under § 510(m) of the act (21 U.S.C. 360(m)) concerning exemptions from premarket notification.

(b) These officials may not further redelegate this authority.

§ 5.402 Detention of adulterated or misbranded medical devices.

(a) The following officials are authorized to perform all the functions of the